

[54] **FERRIC IRON REMOVAL FROM ALUMINUM FLUORIDE SOLUTIONS**

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**Related U.S. Application Data**

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[58] Field of Search ..... **423/112, 126, 132, 139, 423/150, 483, 488, 489, 658.5, 484, 626**

[56] **References Cited**

**U.S. PATENT DOCUMENTS**

2,155,119	4/1939	Ebner .....	423/DIG. 1
2,920,938	1/1960	Matoush .....	423/126
3,145,081	8/1964	Surls et al. ....	423/112
3,211,521	10/1965	George .....	423/139
3,320,032	5/1967	Feller .....	423/139
3,729,541	4/1973	Grunig et al. ....	423/112
3,961,030	6/1976	Wiewiorowski et al. ....	423/132
3,966,909	6/1976	Grunig .....	75/101 BE

**OTHER PUBLICATIONS**

Marcus et al., *Ion Exchange and Solvent Extraction of Metal Complexes*, Wiley Interscience (1969), pp. 523-525, 534-536, 538,546,550.

Peppard et al., "Acid Esters of Orthophosphoric Acid as Selective Extractants for Metallic Cations-Tracer Studies", *J. Inorg. and Nucl. Chem.*, vol. 7 (1958), pp. 269, 276-285.

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[57] **ABSTRACT**

A process for the selective separation of ferric iron from an aluminum fluoride aqueous solution containing ferric iron is effected by contacting said solution with an organic extractant phase consisting essentially of a mixture of a mono (higher alkyl-substituted phenyl) phosphoric acid and a di(higher alkyl-substituted phenyl) phosphoric acid dissolved in an inert diluent to thereby transfer the ferric iron to said organic extractant phase. The efficiency of the process is further improved by adjusting the aluminum fluoride aqueous feed solution to a temperature in the range of about 140° F. to about 175° F. prior to contacting it with the organic extractant phase, maintaining the temperature within this range throughout the ferric iron extraction stage or stages, then heating the resulting iron-depleted aqueous raffinate to a temperature higher than 195° F., and then crystallizing aluminum fluoride trihydrate out of the raffinate at this higher temperature.

**11 Claims, 2 Drawing Figures**

